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EUROPEAN COMMISSION



Brussels, 01-02-2011 C(2011)688

COMMISSION DECISION

of 01-02-2011

amending the marketing authorisation granted by Decision C(2004)3160 for "Ariclaim - duloxetine hydrochloride", a medicinal product for human use

(ONLY THE DUTCH TEXT IS AUTHENTIC)

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amending the marketing authorisation granted by Decision C(2004)3160 for "Ariclaim - duloxetine hydrochloride", a medicinal product for human use

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency¹,

Having regard to Commission Regulation (EC) No 1234/2008 of 24 November 2008 concerning the examination of variations to the terms of marketing authorisations for medicinal products for human use and veterinary medicinal products², and in particular Article 20(7)(a) thereof,

Having regard to the application submitted on 17 October 2010 by Eli Lilly Nederland B.V. under Article 20(3) of Regulation (EC) No 1234/2008,

Having regard to the opinion of the European Medicines Agency, formulated on 16 December 2010 by the Committee for Medicinal Products for Human Use,

Whereas:

- (1) The European Medicines Agency has informed the marketing authorisation holder and the Commission of its favourable opinion on the major variation type II and of the need to amend the marketing authorisation for the medicinal product "Ariclaim duloxetine hydrochloride" which is entered in the Community Register of Medicinal Products under numbers EU/1/04/283/008-012.
- (2) The marketing authorisation should be updated and Decision C(2004)3160 of 11 August 2004 should therefore be amended accordingly. The Community Register of Medicinal Products should also be updated.

HAS ADOPTED THIS DECISION:

Article 1

Decision C(2004)3160 is amended as follows:

OJ L 334, 12.12.2008, p. 7.

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OJ L 136, 30.4.2004, p. 1.

- 1) Annex I is replaced by the text set out in Annex I to this Decision;
- 2) Annex II is replaced by the text set out in Annex II to this Decision;
- 3) Annex III is replaced by the text set out in Annex III to this Decision.

Article 2

This Decision is addressed to Eli Lilly Nederland B.V., Grootslag 1-5, 3991 RA Houten, Nederland.

Done at Brussels, on 01-02-2011.

For the Commission Paola TESTORI COGGI Director-General